108TH CONGRESS 2D SESSION

H. R. 4420

To ensure that women seeking an abortion are fully informed regarding the pain experienced by their unborn child.

IN THE HOUSE OF REPRESENTATIVES

May 20, 2004

Mr. Smith of New Jersey (for himself, Mr. Hyde, Mrs. Jo Ann Davis of Virginia, Mr. Alexander, Mr. Pitts, Mrs. Myrick, Mr. Souder, Mr. Burgess, Mr. Lincoln Diaz-Balart of Florida, Ms. Ros-Lehtinen, Mr. Aderholt, Mr. Tiahrt, Mr. Crane, Mr. Neugebauer, Mr. Franks of Arizona, Mr. Istook, Mr. Akin, Mr. Stearns, Mr. Renzi, Mr. Shimkus, Mr. Pence, Mr. Demint, Mr. Collins, Mr. Brady of Texas, Mr. King of Iowa, Mr. Tancredo, Mr. Ryun of Kansas, and Mr. Toomey) introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

To ensure that women seeking an abortion are fully informed regarding the pain experienced by their unborn child.

- 1 Be it enacted by the Senate and House of Representa-
- 2 tives of the United States of America in Congress assembled,
- 3 SECTION 1. SHORT TITLE.
- 4 This Act may be cited as the "Unborn Child Pain
- 5 Awareness Act of 2004".
- 6 SEC. 2. FINDINGS.
- 7 Congress makes the following findings:

- (1) At least 20 weeks after fertilization, an unborn child has the physical structures necessary to experience pain.
 - (2) There is substantial evidence that by 20 weeks after fertilization, unborn children draw away from certain stimuli in a manner which in an infant or an adult would be interpreted as a response to pain.
 - (3) Anesthesia is routinely administered to unborn children who have developed 20 weeks or more past fertilization who undergo prenatal surgery.
 - (4) There is substantial evidence that the abortion methods most commonly used 20 weeks after fertilization cause substantial pain to an unborn child, whether by dismemberment, poisoning, penetrating or crushing the skull, or other methods. Examples of abortion methods used 20 weeks after fertilization include, but are not limited to the following:
 - (A) The Dilation and Evacuation (DE) method of abortion is commonly performed in the second trimester of pregnancy. In a dilation and evacuation abortion, the unborn child's body parts are grasped at random with a long-toothed clamp. The fetal body parts are then

- torn off of the body and pulled out of the vaginal canal. The remaining body parts are grasped and pulled out until only the head remains. The head is then grasped and crushed in order to remove it from the vaginal canal.
 - (B) Partial-Birth Abortion is an abortion in which the abortion practitioner delivers an unborn child's body until only the head remains inside the womb, punctures the back of the child's skull with a sharp instrument, and sucks the child's brains out before completing the delivery of the dead infant.
 - (5) Expert testimony confirms that by 20 weeks after fertilization an unborn child may experience substantial pain even if the woman herself has received local analysis or general anesthesia.
 - (6) Medical science is capable of reducing such pain through the administration of anesthesia or other pain-reducing drugs directly to the unborn child.
 - (7) There is a valid Federal Government interest in reducing the number of events in which great pain is inflicted on sentient creatures. Examples of this are laws governing the use of laboratory animals and requiring pain-free methods of slaughtering live-

1	stock, which include, but are not limited to the fol-
2	lowing:

- (A) Section 2 of the Humane Slaughter Act (7 U.S.C. 1902) states, "No method of slaughter or handling in connection with slaughtering shall be deemed to comply with the public policy of the United States unless it is humane. Either of the following two methods of slaughtering and handling are hereby found to be humane:
 - "(i) in the case of cattle, calves, horses, mules, sheep, swine, and other live-stock, all animals are rendered insensible to pain by a single blow or gunshot or an electrical, chemical or other means that is rapid and effective, before being shackled, hoisted, thrown, cast, or cut; or
 - "(ii) by slaughtering in accordance with the ritual requirements of the Jewish faith or any other religious faith that prescribes a method of slaughter whereby the animal suffers loss of consciousness by anemia of the brain caused by the simultaneous and instantaneous severance of the carotid arteries with a sharp instrument

1	and handling in connection with such
2	slaughtering.".
3	(B) Section 13(a)(3) of the Animal Wel-
4	fare Act (7 U.S.C. 2143(a)(3)) sets the stand-
5	ards and certification process for the humane
6	handling, care, treatment, and transportation of
7	animals. This includes having standards with
8	respect to animals in research facilities that in-
9	clude requirements—
10	"(i) for animal care, treatment, and
11	practices in experimental procedures to en-
12	sure that animal pain and distress are
13	minimized, including adequate veterinary
14	care with the appropriate use of anesthetic,
15	analgesic, tranquilizing drugs, or eutha-
16	nasia;
17	"(ii) that the principal investigator
18	considers alternatives to any procedure
19	likely to produce pain to or distress in an
20	experimental animal;
21	"(iii) in any practice which could
22	cause pain to animals—
23	"(I) that a doctor of veterinary
24	medicine is consulted in the planning
25	of such procedures;

1	"(II) for the use of tranquilizers,
2	analgesics, and anesthetics;
3	"(III) for pre-surgical and post-
4	surgical care by laboratory workers, in
5	accordance with established veterinary
6	medical and nursing procedures;
7	"(IV) against the use of para-
8	lytics without anesthesia; and
9	"(V) that the withholding of
10	tranquilizers, anesthesia, analgesia, or
11	euthanasia when scientifically nec-
12	essary shall continue for only the nec-
13	essary period of time;".
14	(C) Section 495 of the Public Health Serv-
15	ice Act (42 U.S.C. 289d) directs the Secretary
16	of Health and Human Services, acting through
17	the Director of the National Institutes of
18	Health, to establish guidelines for research fa-
19	cilities as to the proper care and treatment of
20	animals, including the appropriate use of tran-
21	quilizers, analgesics, and other drugs, except
22	that such guidelines may not prescribe methods
23	of research. Entities that conduct biomedical
24	and behavioral research with National Insti-
25	tutes of Health funds must establish animal

1	care committees which must conduct reviews at
2	least semi-annually and report to the Director
3	of such Institutes at least annually. If the Di-
4	rector determines that an entity has not been
5	following the guidelines, the Director must give
6	the entity an opportunity to take corrective ac-
7	tion, and, if the entity does not, the Director
8	must suspend or revoke the grant or contract
9	involved.
10	SEC. 3. AMENDMENT TO THE PUBLIC HEALTH SERVICE
11	ACT.
12	The Public Health Service Act (42 U.S.C. 201 et
13	seq.) is amended by adding at the end the following:
14	"TITLE XXIX—UNBORN CHILD
15	PAIN AWARENESS
16	"SEC. 2901. DEFINITIONS.
17	"In this title:
18	"(1) Abortion.—The term 'abortion' means
19	the intentional use or prescription of any instru-

"(1) ABORTION.—The term 'abortion' means the intentional use or prescription of any instrument, medicine, drug, or any other substance or device to terminate the pregnancy of a woman known to be pregnant with an intention other than to increase the probability of a live birth, to preserve the life or health of the child after live birth, or to remove a dead fetus.

1 "(2) ABORTION PROVIDER.—The term 'abortion 2 provider' means any person legally qualified to per-3 form an abortion under applicable Federal and State 4 laws.

"(3) Pain-capable unborn child.—

- "(A) IN GENERAL.—The term 'pain-capable unborn child' means an unborn child who has reached a probable stage of development of 20 weeks after fertilization.
- "(B) RULE OF CONSTRUCTION.—Nothing in subparagraph (A) shall be construed as a determination or finding by Congress that pain may not in fact be experienced by an unborn child at stages of development prior to 20 weeks after fertilization.
- "(4) Probable age of development' means the duration of development after fertilization of the unborn child at the time an abortion is performed, as determined in the good faith judgment of the abortion provider on the basis of examination of the unborn child using ultrasound or other imaging technology, in addition to information obtained by interviewing the pregnant woman.

1	"(5) Unborn Child.—The term 'unborn child'
2	means a member of the species homo sapiens, at any
3	stage of development, who is carried in the womb.
4	"(6) Woman.—The term 'woman' means a fe-
5	male human being who is capable of becoming preg-
6	nant, whether or not she has reached the age of ma-
7	jority.
8	"SEC. 2902. REQUIREMENT OF INFORMED CONSENT.
9	"(a) Requirement of Compliance by Pro-
10	VIDERS.—An abortion provider performing any abortion,
11	of a pain-capable unborn child, that is in or affecting
12	interstate commerce shall comply with the requirements
13	of this title.
14	"(b) Provision of Consent.—
15	"(1) IN GENERAL.—Before any part of an abor-
16	tion involving a pain-capable unborn child begins,
17	the abortion provider or his or her agent shall pro-
18	vide the pregnant woman involved, by telephone or
19	in person, with the information described in para-
20	graph (2).
21	"(2) Required information.—
22	"(A) Oral statement.—
23	"(i) In general.—An abortion pro-
24	vider or the provider's agent to whom
25	paragraph (1) applies shall make the fol-

1 lowing oral statement to the pregnant 2 woman (or in the case of a deaf or non-3 English speaking woman, provide the statement in a manner that she can easily understand): 6 You are considering having an abortion of 7 an unborn child who will have developed, 8 at the time of the abortion, approximately 9 weeks after fertilization. The Con-10 gress of the United States has determined 11 that at this stage of development, an un-12 born child has the physical structures nec-13 essary to experience pain. There is sub-14 stantial evidence that by this point, unborn 15 children draw away from surgical instru-16 ments in a manner which in an infant or 17 an adult would be interpreted as a re-18 sponse to pain. Congress finds that there 19 is substantial evidence that the process of 20 being killed in an abortion will cause the 21 unborn child pain, even though you receive 22 a pain-reducing drug or drugs. Under the 23 Federal Unborn Child Pain Awareness Act 24 of 2004, you have the option of choosing to 25 have anesthesia or other pain-reducing 1 drug or drugs administered directly to the 2 pain-capable unborn child if you so desire. The purpose of administering such drug or 3 drugs would be to reduce or eliminate the capacity of the unborn child to experience 6 pain during the abortion procedure. In 7 some cases, there may be some additional 8 risk to you associated with administering 9 such a drug.'.

"(ii) Description of Risks.—After making the statement required under clause (i), the abortion provider may provide the woman involved with his or her best medical judgment on the risks of administering such anesthesia or analgesic, if any, and the costs associated therewith.

"(iii) Administration of anes-Thesia.—If the abortion provider is not qualified or willing to administer the anesthesia or other pain-reducing drug in response to the request of a pregnant women after making the statement required under clause (i), the provider shall—

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1 "()	I) arrange for a qualified spe-
2 cialist	to administer such anesthesia
3 or drug	; or
4 "()	II) advise the pregnant
5 woman-	
6	"(aa) where she may obtain
7 suc	ch anesthesia or other pain re-
8 du	cing drugs for the unborn child
9 in	the course of an abortion; or
10	"(bb) that the abortion pro-
11 vid	ler is unable to perform the
12 ab	ortion if the woman elects to
13 rec	ceive anesthesia or other pain-
14 rec	ducing drug for her unborn
15 chi	ild.
16 "(iv)	Rule of construction.—
Nothing in	this section may be construed
to impede	an abortion provider or the
19 abortion pr	ovider's agent from offering
their own ev	valuation on the capacity of the
21 unborn child	d to experience pain, the advis-
ability of	administering pain-reducing
drugs to th	e unborn child, or any other
matter, as l	ong as such provider or agent
provides the	e required information, obtains

the woman's signature on the decision form, and otherwise complies with the affirmative requirements of the law.

"(B) Unborn Child Pain Awareness
Brochure.—An abortion provider to whom
paragraph (1) applies shall provide the pregnant woman with the Unborn Child Pain
Awareness Brochure (referred to in this section
as the 'Brochure') to be developed by the Department of Health and Human Services under
subsection (c) or with the information described
in subsection (c)(2) relating to accessing such
Brochure.

"(C) Unborn Child Pain Awareness
Decision form.—An abortion provider to
which paragraph (1) applies shall provide the
pregnant woman with the Unborn Child Pain
Awareness Decision Form (provided for under
subsection (c)) and obtain the appropriate signature of the woman on such form.

21 "(c) Unborn Child Pain Awareness Bro-22 Chure.—

"(1) Development.—Not later than 90 days
after the date of enactment of this title, the Secretary shall develop an Unborn Child Pain Aware-

ness Brochure. Such Brochure shall be written in English and Spanish and shall contain the same information as required under the statement under subsection (b)(2)(A)(i), including greater detail on her option of having a pain-reducing drug or drugs administered to the unborn child to reduce the experience of pain by the unborn child during the abortion. Such information shall be written in an objective and nonjudgmental manner and be printed in a typeface large enough to be clearly legible. The Brochure shall be made available by the Secretary at no cost to any abortion provider.

- "(2) Internet information.—The Brochure under this section shall be available on the Internet website of the Department of Health and Human Services at a minimum resolution of 70 DPI (dots per inch). All pictures appearing on the website shall be a minimum of 200x300 pixels. All letters on the website shall be a minimum of 12 point font. All such information and pictures shall be accessible with an industry standard browser, requiring no additional plug-ins.
- "(3) Presentation of Brochure.—An abortion provider or his or her agent must offer to provide a pregnant woman with the Brochure, developed

1	under paragraph (1), before any part of an abortion
2	of a pain-capable child begins—
3	"(A) through an in-person visit by the
4	pregnant woman;
5	"(B) through an e-mail attachment, from
6	the abortion provider or his or her agent; or
7	"(C) through a request to have such Bro-
8	chure mailed, by certified mail, to the woman at
9	least 72 hours before any part of the abortion
10	begins.
11	"(4) Waiver.—After the abortion provider or
12	his or her agent offers to provide a pregnant woman
13	the Brochure, a pregnant woman may waive receipt
14	of the Brochure under this subsection by signing the
15	waiver form contained in the Unborn Child Pain
16	Awareness Decision Form.
17	"(5) Unborn Child Pain Awareness Deci-
18	SION FORM.—Not later than 30 days after the date
19	of enactment of this title, the Secretary shall develop
20	an Unborn Child Pain Awareness Decision Form.
21	To be valid, such Form shall—
22	"(A) with respect to the pregnant
23	woman—
24	"(i) contain a statement that affirms
25	that the woman has received or been of-

1	fered all of the information required in
2	subsection (b);
3	"(ii) require the woman to explicitly
4	either request or refuse the administration
5	of pain-reducing drugs to the unborn child;
6	"(iii) be signed by a pregnant woman
7	prior to the performance of an abortion in-
8	volving a pain-capable unborn child; and
9	"(B) with respect to the abortion pro-
10	vider—
11	"(i) contain a statement that the pro-
12	vider has provided the woman with all of
13	the information required under subsection
14	(b);
15	"(ii) if applicable, contain a certifi-
16	cation by the provider that an exception
17	described in section 2903 applies and the
18	detailed reasons for such certification; and
19	"(iii) be signed by the provider prior
20	to the performance of the abortion proce-
21	dure.
22	"(6) Maintenance of Records.—The Sec-
23	retary shall promulgate regulations relating to the
24	period of time during which copies of Forms under

1 paragraph (5) shall be maintained by abortion pro-2 viders. 3 "SEC. 2903. EXCEPTION FOR MEDICAL EMERGENCIES. 4 "(a) In General.—The provisions of section 2902 shall not apply to an abortion provider in the case of a 6 medical emergency. 7 "(b) Medical Emergency Defined.— "(1) IN GENERAL.—In subsection (a), the term 8 9 'medical emergency' means a condition which, in the 10 reasonable medical judgment of the abortion pro-11 vider, so complicates the medical condition of the 12 pregnant woman that a delay in commencing an 13 abortion procedure would impose a serious risk of 14 causing grave and irreversible physical health dam-15 age entailing substantial impairment of a major bod-16 ily function. 17 "(2) Reasonable medical judgment.—In 18 paragraph (1), the term 'reasonable medical judg-19 ment' means a medical judgment that would be 20 made by a reasonably prudent physician, knowledge-21 able about the case and the treatment possibilities 22 with respect to the medical conditions involved. 23 "(c) Certification.— "(1) IN GENERAL.—Upon a determination by 24 25 an abortion provider under subsection (a) that a

- 1 medical emergency exists with respect to a pregnant
- woman, such provider shall certify the specific med-
- 3 ical conditions that constitute the emergency.
- 4 "(2) False statements.—An abortion pro-
- 5 vider who willfully falsifies a certification under
- 6 paragraph (1) shall be subject to all the penalties
- 7 provided for under section 2904 for failure to com-
- 8 ply with this title.

9 "SEC. 2904. PENALTIES FOR FAILURE TO COMPLY.

- 10 "(a) IN GENERAL.—An abortion provider who will-
- 11 fully fails to comply with the provisions of this title shall
- 12 be subject to civil penalties in accordance with this section
- 13 in an appropriate Federal court.
- 14 "(b) Commencement of Action.—The Attorney
- 15 General, the Deputy Attorney General, the Associate At-
- 16 torney General, or any Assistant Attorney General or
- 17 United States Attorney who has been specifically des-
- 18 ignated by the Attorney General may commence a civil ac-
- 19 tion under this section.
- 20 "(c) Certification Requirements.—At the time
- 21 of the commencement of an action under this section, the
- 22 Attorney General, the Deputy Attorney General, the Asso-
- 23 ciate Attorney General, or any Assistant Attorney General
- 24 or United States Attorney who has been specifically des-
- 25 ignated by the Attorney General to commence a civil ac-

- 1 tion under this section, shall certify to the court involved
- 2 that, at least 30 calendar days prior to the filing of such
- 3 action, the Attorney General, the Deputy Attorney Gen-
- 4 eral, the Associate Attorney General, or any Assistant At-
- 5 torney General or United States Attorney involved—
- 6 "(1) has provided notice of the alleged violation
- 7 of this section, in writing, to the Governor or Chief
- 8 Executive Officer and Attorney General or Chief
- 9 Legal Officer of the State or political subdivision in-
- volved, as well as to the State medical licensing
- board or other appropriate State agency; and
- "(2) believes that such an action by the United
- 13 States is in the public interest and necessary to se-
- cure substantial justice.
- 15 "(d) First Offense.—Upon a finding by a court
- 16 that a respondent in an action commenced under this sec-
- 17 tion has knowingly violated a provision of this title, the
- 18 court shall notify the appropriate State medical licensing
- 19 authority in order to effect the suspension of the respond-
- 20 ent's medical license in accordance with the regulations
- 21 and procedures promulgated under section 2905, or shall
- 22 assess a civil penalty against the respondent in an amount
- 23 not to exceed \$100,000, or both.
- "(e) SECOND OFFENSE.—Upon a finding by a court
- 25 that the respondent in an action commenced under this

- 1 section has knowingly violated a provision of this title and
- 2 the respondent has been found to have knowingly violated
- 3 a provision of this title on a prior occasion, the court shall
- 4 notify the appropriate State medical licensing authority in
- 5 order to effect the revocation of the respondent's medical
- 6 license in accordance with the regulations and procedures
- 7 promulgated under section 2905, or shall assess a civil
- 8 penalty against the respondent in an amount not to exceed
- 9 \$250,000, or both.
- 10 "(f) Hearing.—With respect to an action under this
- 11 section, the appropriate State medical licensing authority
- 12 shall be given notification of and an opportunity to be
- 13 heard at a hearing to determine the penalty to be imposed
- 14 under this section.
- 15 "(g) Private Right of Action.—A pregnant
- 16 woman upon whom an abortion has been performed in vio-
- 17 lation of this title, or the parent or legal guardian of such
- 18 a woman if she is an unemancipated minor, may com-
- 19 mence a civil action against the abortion provider for any
- 20 knowing or reckless violation of this title for actual and
- 21 punitive damages.
- 22 "SEC. 2905. REGULATIONS.
- 23 "A State, and the medical licensing authority of the
- 24 State, shall promulgate regulations and procedures for the
- 25 revocation or suspension of the medical license of an abor-

- 1 tion provider upon a finding by a court under section 2904
- 2 that the provider has violated a provision of this title. A
- 3 State that fails to implement such procedures shall be sub-
- 4 ject to loss of funding under title XIX of the Social Secu-
- 5 rity Act (42 U.S.C. 1396 et seq.).".

6 SEC. 4. PREEMPTION.

- 7 Nothing in this Act or the amendments made by this
- 8 Act shall be construed to preempt any provision of State
- 9 law to the extent that such State law establishes, imple-
- 10 ments, or continues in effect greater protections for un-
- 11 born children from pain than the protections provided
- 12 under this Act and the amendments made by this Act.

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